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UTILITY PATENT APPLICATION TRANSMITTAL

(Only for new nonprovisional applications under 37 CFR 1.53(b))

Attorney Docket No.

503.34465VC3

Total Pages

First Named Inventor or Application Identifier

HONBO, et al.

Express Mail Label No.

APPLICATION ELEMENTS

See MPEP chapter 600 concerning utility patent application contents.

ADDRESS TO:

Assistant Commissioner for Patents
Box Patent Application
Washington, DC 20231

1. ☒ Fee: \$1686.00

Please charge any shortages in the fees or credit any overpayments thereof to the deposit account of Antonelli, Terry, Stout & Kraus, Deposit Account No. 01-2135.

6. ☐ Microfiche Computer Program (Appendix)

7. Nucleotide and/or Amino Acid Sequence Submission (if applicable, all necessary)

- a. ☐ Computer Readable Copy
b. ☐ Paper Copy (identical to computer copy)
c. ☐ Statement verifying identity of above copies

2. ☒ Specification Total Pages 38

3. ☒ Drawing(s) (35 USC 113) Total Sheets 5

4. Oath or Declaration Total Pages 2

- a. ☐ Newly executed (original or copy)
b. ☒ Copy from a prior application (37 CFR 1.63(d)) (for continuation/divisional with Box 17 completed) [Note Box 5 below]

i. ☐ DELETION OF INVENTOR(S)
Signed statement attached deleting inventor(s) named in the prior application, see 37 CFR 1.63(d)(2) and 1.33(b).

5. ☒ Incorporation By Reference (useable if Box 4b is checked)
The entire disclosure of the prior application, from which a copy of the oath or declaration is supplied under Box 4b, is considered as being part of the disclosure of the accompanying application and is hereby incorporated by reference therein.

ACCOMPANYING APPLICATION PARTS

8. ☐ Assignment Papers (cover sheet & document(s))
9. ☐ 37 CFR 3.73(b) Statement (when there is an assignee) ☐ Power of Attorney
10. ☐ English Translation Document (if applicable)
11. ☒ Information Disclosure Statement (IDS)/PTO-1449 ☐ Copies of IDS Citations
12. ☒ Preliminary Amendment
13. ☒ Return Receipt Postcard (MPEP 503) (Should be specifically itemized)
14. ☐ Small Entity ☐ Statement filed in prior application, Status still proper and desired
15. ☐ Certified Copy of Priority Document(s) (if foreign priority is claimed)
16. ☒ Other: Claim for Priority

17. If a CONTINUING APPLICATION, check appropriate box and supply the requisite information:

☐ Continuation ☒ Divisional ☐ Continuation-in-part (CIP) of prior application No: 08/630,501

18. CORRESPONDENCE ADDRESS

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11. SIGNATURE OF APPLICANT, ATTORNEY, OR AGENT REQUIRED

NAME William I. Solomon

SIGNATURE *William I. Solomon*

DATE December 28, 1999

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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant(s): HONBO, et al.
Serial No.: Divisional of application
Serial No. 08/630,501
Filed: December 28, 1999
For: NON-AQUEOUS SECONDARY BATTERY
Group:
Examiner:

PRELIMINARY AMENDMENT

Assistant Commissioner for Patents
Washington, D.C. 20231

December 28, 1999

Sir:

Please amend the above-identified application, prior to
examination thereof, as follows:

IN THE SPECIFICATION

Please amend the specification as follows:

Page 1, lines 4 and 5, in the title of the invention,
delete "NON-AQUEOUS SECONDARY BATTERY AND A MANUFACTURING
METHOD OF GRAPHITE POWDER" and insert --NON-AQUEOUS SECONDARY
BATTERY AND A METHOD OF MANUFACTURING GRAPHITE POWDER--;

line 6, insert the following paragraph:

--This application is a Divisional application of
application Serial No. 08/630,501, filed April 10, 1996.--;

line 7, delete "(1) Field of the Invention:";

line 13, after "uses" delete "the";

line 15, after "for" delete "using" and insert

--use--;

line 17, delete "(2) Description of the Prior Art:";

line 19, after "problems" delete "in" and insert --relating to--;

line 21, after "during" delete "repeating charge" and insert --repeated charging-- and after "and" delete "discharge" and insert --discharging--;

line 22, after "lithium" delete "will grow" and insert --grows--;

line 23, after "circuit" delete "is" and insert --will be--;

line 25, after "material" insert --is disclosed-- and insert --has been proposed--; and

line 26, after "material" delete "replaceable" and insert --in place--; and

line 27, after "reactions" delete "are" and insert --involve-- and after "lithium" delete "ions" and insert --ion--.

Page 2, line 2, after "and" insert --so-- and after "hardly" delete "deposit" and insert --deposited--;

line 7, after "lithium." delete "The" and insert --Thus,--;

line 9, after "use" delete "the above";

line 11, after "increase" insert --the--;

line 12, after "which" delete "is" and insert --constitutes--;

line 13, delete "proceed" and insert --allow--
and after "reactions" insert --to proceed--;

line 16, after "as" delete "it", after "from"
delete "a" and insert --the-- and after "that" delete "the";

line 21, after "effects" insert --of the
pulverizing process--;

line 23, after "when" delete "the above";

line 25, after "disadvantage" delete "is
caused" and insert --results in--; and

line 27, after "rapid" delete "charge and
discharge" and insert --charging and discharging--.

Page 3, line 5, delete "(1) Objects of the Invention:";

line 7, after "problems" delete "involved in
the prior arts" and after "to" delete "disclose" and insert
--provide--;

line 11, after "in" delete "the" and insert
--its--;

line 12, after "above" delete "disclosed" and
insert --mentioned--;

line 13, delete "(2) Methods of solving the
Problems:";

line 23, after "in" delete "the"; and

line 24, after "diffraction" insert --of the material--.

Page 4, line 4, after "powder" insert --to--;
line 5, after "powder" insert --to--;
line 7, after "heated" insert --to--;
line 8, after "of" insert --the--;
line 19, after "group" delete "consisted" and insert --consisting--; and
line 21, after "treatment" insert a comma --,--

Page 5, line 8, after "in" (first occurrence) delete "a" and insert --the--, after "of" insert --a-- and after "to" delete "proceed" and insert --facilitate--;
line 10, after "which" delete "is" and insert --constitutes--;
line 21, after "However," insert --for--;
line 24, after "graphite" delete "powder" and insert --powders--; and
line 27, after "various" delete "time" and insert --times-- and after "And," insert --the--.

Page 6, line 1, after "graphite" delete "powder" and insert --powders--;

line 2, delete "searching" and insert
--determining--;
line 4, after "had" delete "the";
line 11, after "graphite" insert --samples--;
line 16, after "of" insert --the-- and after
"and" (second occurrence) insert --the--;
line 19, after "of" insert --the-- and after
"and" insert --the--; and
line 20, after "of" (first occurrence) insert
--the--.

Page 7, line 3, after "of" (second occurrence) insert
--the--;
line 4, after "As" delete "the" and insert
--a-- and after "that" delete "the";
line 17, after "of" insert --the--;
line 19, delete "the" and insert --a-- and
after "that" delete "the";
line 26, after "for" delete "existing" and
insert --existence of--; and
line 27, after "assumed" insert --to be--.

Page 8, line 1, after "having" delete "the" and insert
--a--;

line 2, after "to" delete "the" and after
"having" delete "the";

line 4, after "shocks." delete "The graphite"
and insert --Graphite-- and after "powder" delete "having" and
insert --of--;

line 12, after "assumed" insert --to be due to
the fact--;

line 13, after "has" insert --been--;

line 14, after "and" (first occurrence) insert
--has-- and after "carbon," delete "and" (second occurrence);"

line 15, delete "proceeding of" and insert
--with the result that--;

line 18, after "Analysis" delete "on" and
insert --of--;

line 19, after "that" delete "the";

line 20, delete "contained" and insert
--present in an amount--;

line 21, after "from" insert --a--;

line 22, after "apparatus" insert a comma
--,--;

line 23, after "like" insert a comma --,-- and
after "graphite" delete "at" and insert --during--; and

line 25, after "capacity" insert a comma --,--.

Page 9, line 1, after "invention," delete "the" and
insert --a--;

line 7, after "to" insert --an amount--;

line 12, after "for" delete "the" and insert
--a--;
line 14, delete "usages" and insert
--purposes--;
line 15, before "equal" insert --a size--;
line 18, after "same" delete "is" and insert
--will be--;
line 20, after "obtaining" delete "the";
line 26, after "the" delete "above"; and
line 17, after "one" delete "of which" and
insert --having a--.

Page 10, line 1, after "line" delete "is appeared" and
insert --which appears--;
line 2, after "range" delete "form" and insert
--from--;
line 3, after "is," insert --where--;
line 5, after "desirable." delete "Because" and
insert --As a result-- and before "graphite" delete "the" and
insert --a--;
line 9, after "to" (first occurrence) delete
"the" and insert --a--;
line 11, after "than" insert --in--;
line 13, after "contains" delete "the";
line 14, after "having" delete "the" and insert
--a--;
line 17, after "is" delete "made decrease" and
insert --decreased--;

line 19, after "performed" insert --to--; and
line 26, after "to" (first occurrence) delete
"the" and insert --a--.

Page 11, line 5, after "pulverized," delete "the" and
after "containing" delete "the" and insert --a--;

line 6, after "structure" insert --in a
quantity--;

line 9, after "graphite" insert --in a
quantity-- and after "by" insert --employing--;

line 12, after "invention" (second occurrence)
insert --proposes--;

line 14, after "particularly" delete "much" and
insert --a large amount--;

line 15, after "materials" insert --to--;

line 20, after "is" delete "as" and insert
--the--; and

line 25, after "group" delete "consisted" and
insert --consisting--.

Page 12, line 11, after "by" delete "an";

line 12, after "of" delete "the";

line 13, after "of" delete "the" and after
"less" insert --than--;

line 14, delete "the";

line 17, after "decreased" insert --to--;

line 23, after "and" insert --the--; and

line 25, after "As" delete "the" and insert
--a--.

Page 13, line 8, insert --Where the fraction of the
rhombohedral structure is 10% or less, and from Equations (1)
and (2) herein, respectively, P_2/P_1 is 0.41 or less and P_3/P_1 is
0.33 or less.-- after "increased";

line 10, after "hardly" insert --will--;

line 23, after "As" delete "the" and insert
--a-- and after "evaluation" delete "on" and insert --of--;
and

line 27, after "and" (second occurrence) insert
--a--.

Page 14, line 4, after "assumed" insert --to relate to
the fact--;

line 9, after "impurities" insert a comma
--,--;

line 15, after "the" (second occurrence) delete
"charge" and insert --charging--;

line 16, after "and" delete "discharge" and
insert --discharging--;

line 19, after "group" delete "consisted" and
insert --consisting--;"

line 21, after "of" insert --the--;

line 22, after "group" delete "consisted" and
insert --consisting--;

line 25, after "because" delete ", an" and
insert --the--; and

line 26, after "solution" delete "is" and
insert --will be--.

Page 15, line 2, after "1" insert --is a graph which--
and after "of" delete "the";

line 3, after "graphite" delete the comma ","
and insert a semicolon --;--;

line 4, after "2" insert --is a graph which--
and after "of" delete "the";

line 6, after "C)" delete the comma "," and
insert a semicolon --;--;

line 7, after "3" insert --is a graph which--;

line 9, after "C)" delete the comma "," and
insert a semicolon --;--;

line 10, after "4" insert --is a graph which--;

line 11, after "1" delete the comma "," and
insert a semicolon --;--;

line 12, after "5" insert --is a graph which--;

line 14, after "invention" delete the comma ","
and insert a semicolon --;--;

line 15, after "6" delete "indicates" and
insert --is--;

line 17, after "2" delete the comma "," and
insert a semicolon --;--;

line 19, delete "figure" and insert
--Figure--;"

line 20, after "capacity" delete the comma ","
and insert a semicolon --;--;

line 23, after "invention" delete the comma ","
and insert a semicolon --;--;

line 25, after "of" delete "repeating" and
insert --repetitions of--; and

line 26, after "cycles" delete the comma ","
and insert --; and--.

Page 16, line 2, after "current" delete the comma ","
and insert a period --.--;

line 6, after "to" insert --the--;

line 7, after "invention" delete "are" and
insert --will be--;

line 12, after "to" delete "be" and insert
--form a-- and after "which" insert --the--;

line 13, before "a" delete "by" and insert
--using--;

line 14, after "obtain" insert --a--; and

line 24, after "by" delete "an".

Page 17, line 1, after "voltage" delete the semicolon
";" and insert --of-- and after "current" delete the semicolon
";" and insert --of--;

line 2, after "source" delete the semicolon ";"
and insert --of--;

line 20, after "to" delete "compare" and insert
--provide a comparison--;

line 21, after "invention," delete "the";

line 22, after "obtain" delete "the";

line 26, after "belong" delete "with" and
insert --to--; and

line 27, after "means" delete the comma "," and
insert --that--.

Page 18, line 4, after "2," delete "the";

line 7, after "acid" insert --comprised--;

line 15, after "belong" delete "with" and insert
--to--;

line 23, after "words," delete "a" and insert
--the--; and

line 25, after "battery" insert a comma --,--.

Page 19, line 3, after "pyrolidone" insert a comma
--,--, after "which" insert --the-- and after "heating" delete
"tempearture" and insert --temperatures--;

line 5, before "and" delete "thick," and insert
--thickness--;

line 6, after "in" insert --a--;

line 7, after "which" insert --the--;

line 8, after "range" insert --of--;

line 10, after "foil" delete "applied";

line 11, after "the" insert --applied-- and
after "mm" insert --sheet--;
line 17, after "metal" insert --element--;
line 18, after "electrode," insert --and--;
line 20, after "screwed" insert --on--;
line 23, after "diethylcarbonate" delete "by"
and insert --at a ratio of--; and
line 24, after "with" insert --a--.

Page 20, line 7, after "to" delete "compare" and insert
--provide a comparison--;
line 19, after "powder" delete "is" and insert
--will be--;
line 22, after "and" delete "an" and insert
--the--; and
line 26, after "which" insert --the-- and after
"temperature" delete "at" and insert --during--.

Page 21, line 2, after "which" insert --the-- and after
"temperature" delete "at" and insert --during--; and
line 19, after "means" delete the comma "," and
insert --that--.

Page 22, line 5, after "in" delete "accordance with
the";
line 6, after "substantially" insert --the--;
and

line 12, after "as" (first occurrence) delete
"same as" and insert --in--.

Page 24, line 4, after "particular," delete "the" and
insert --an--;

line 10, after "member" delete "assigned with"
and insert --identified by--; and

line 11, after "the" insert --reference-- and
after "numeral" delete "mark".

Page 25, line 2, after " μ m" delete "thick" and insert
--thickness--;

line 3, after "in" insert --a--;

line 4, after " μ m" delete "thick" and insert
--thickness--;

line 7, after "foil" delete "applied" and after
"the" (second occurrence) insert --applied--;

line 9, after "sheet" delete "of";

line 12, after "bared" delete "portion" and
insert --portions--;

line 15, after " μ m" delete "thick" and insert
--thickness--;

line 16, after "in" insert --a--;

line 17, after " μ m" delete "thick" and insert
--thickness--;

line 20, after "foil" delete "applied" and
after "the" (second occurrence) insert --applied--;

line 21, after "sheet" delete "of";

line 22, after "as" (first occurrence) delete
"same as" and insert --with--;
line 23, delete "of" (first occurrence); and
line 26, delete "portion" and insert
--portions--.

Page 26, line 1, after " μm " delete "thick" and insert
--thickness--;
line 3, after "negative" insert --electrode--;
line 5, after "of" delete "the";
line 12, after "filled" insert --in--;
line 17, after "steps," delete "the charge and
discharge" and insert --charging and discharging--;
line 18, after "condition" delete "that" and
insert --in which--;
line 19, after "respective" insert --ones--;
line 20, delete "charge" and insert
--charging-- and after "the" delete "discharge was" and insert
--discharging were--;
line 21, after "discharging" delete "current
was" and insert --currents were--;
line 22, after "and" (first occurrence) delete
"the" and after "rapid" delete "charge" and insert
--charging--;
line 23, delete "discharge" and insert
--discharging--;
line 26, after "to" delete "compare" and insert
--provide a comparison--; and

line 27, after "the" insert --same--.

Page 27, line 1, delete "same as the";

line 3, after "characteristics" delete "was"
and insert --were--;

line 4, after "determined" delete "as" and
insert --in the--, after "same" insert --way-- and after "as"
delete "the";

line 5, after "comparison" delete "on" and
insert --of--;

line 7, after "art)" delete "is" and insert
--will be--;

line 9, after "indicates" insert --a--;

line 10, after "the" (first occurrence) delete
"charge and discharge" and insert --charging and
discharging--;

line 15, after "and" delete "a" and insert
--the--;

line 17, after "and" delete "a" and insert
--the--;

line 20, after "charging" insert --current--;

line 21, after "when" delete "the";

line 22, after "rapid" delete "charge" and
insert --charging-- and after "rapid" delete "discharge" and
insert --discharging--; and

line 25, after "With" delete "the" and insert
--a-- and after "charging" insert --current--.

Page 28, line 1, after "of" delete "decreasing" and
insert --decrease of--;

line 2, after "respective" insert --ones--;

line 3, after "of" (first occurrence) insert
--the--;

line 4, after "Therefore," delete "in";

line 5, delete "accordance with" and insert
--by--;

line 7, after "the" (first occurrence) delete
"decreasing" and after "of" insert --decrease of--;

line 10, after "characteristics" delete "in"
and insert --for-- and after "rapid" delete "charge and
discharge" and insert --charging and discharging--;

line 17, after "foil" delete "by as" and insert
--in the--, after "same" insert --manner-- and after "as"
(second occurrence) delete "the";

line 21, after "foil" delete "applied" and
after "the" (second occurrence) insert --applied--;

line 22, after "sheet" delete "of";

line 23, after "portions" delete "of"; and

line 25, after "was" delete "as" and insert
--the--.

Page 29, line 1, after "5" insert a comma --,-- and
after "as" (second occurrence) insert --by--;

line 3, after "welding" insert --a--;

line 6, after "battery," delete "charge and
discharge" and insert --charging and discharging--;

line 7, after "of" (first occurrence) insert
--a--;

line 8, after "the" (second occurrence) delete
"charge and discharge" and insert --charging and
discharging--;

line 9, after "As" delete "the" and insert
--a--;

line 10, after "and" delete "a" and insert
--the--, and after "ratio" delete "in" and insert --of--;

line 14, after "embodiment" delete "is" and
insert --are--;

line 17, after "and" (first occurrence) delete
"an"; and

line 20, after "which" insert --the--.

REMARKS

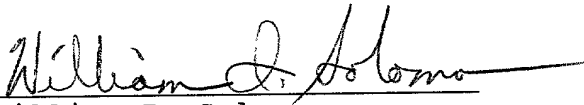
Applicants have amended their specification in light of the history of prior application Serial No. 08/630,501, filed April 10, 1996 (including amendments made to the specification therein in the Preliminary Amendment filed June 28, 1996, and Amendment filed December 12, 1997), and in order to refer to prior application Serial No. 08/630,501 so as to satisfy the requirements of 35 USC 120. It is respectfully submitted that these amendments to the specification do not add new matter to the application.

In view of the foregoing, entry of the present amendments, and examination of the above-identified application in due course, are respectfully requested.

To the extent necessary, Applicants petition for an extension of time under 37 CFR § 1.136. Please charge any shortage in fees due in connection with the filing of this paper, including extension of time fees, to the Deposit Account No. 01-2135 (Case No. 503.34465VC3) and please credit any excess fees to such Deposit Account.

Respectfully submitted,

ANTONELLI, TERRY, STOUT & KRAUS, LLP



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Specification

Title of the Invention

NON-AQUEOUS SECONDARY BATTERY AND A MANUFACTURING
5 METHOD OF GRAPHITE POWDER

Background of the Invention

(1) Field of the Invention:

The present invention relates to a carbon material
10 which intercalates into or deintercalates from lithium, and to
a method for manufacturing the same. In particular, the
present invention relates to a lithium secondary battery,
which uses the carbon material as a negative electrode active
material, having a high energy density and a long life. The
15 lithium battery is suitable for using in portable apparatus,
electric automobiles, power storage, etc.

(2) Description of the Prior Art:

The Lithium secondary battery using lithium metal for the
negative electrode has some problems in safety. For example,
20 lithium easily deposits like dendrite on the lithium metal
negative electrode during repeating charge and discharge of
the battery, and if the dendritic lithium will grow to a
positive electrode, an internal short circuit is caused
between the positive electrode and the negative electrode.

25 Therefore, a carbon material is disclosed as the
negative electrode active material replaceable of lithium
metal. Charge and discharge reactions are lithium ions

intercalation into the carbon material and deintercalation from the carbon material, and lithium is hardly deposit like dendrite. As for the carbon material, graphite is disclosed in JP-B-62-23433 (1987).

5 The graphite disclosed in JP-B-62-23433 (1987) forms an intercalation compound with lithium, because of intercalation or deintercalation of lithium. The graphite is used as a material for the negative electrode of the lithium secondary battery. In order to use the above graphite as the
10 negative active material, it is necessary to pulverize the graphite to increase surface area of the active material, which is a charge and discharge reaction field, so as to proceed the charging and discharging reactions smoothly. Desirably, it is necessary to pulverize the graphite to
15 powder having a particle diameter equal to or less than 100 μm . However, as it is apparent from a fact that the graphite is used as a lubricating material, the graphite easily transfers its layers. Therefore, its crystal structure is changed by the pulverizing process, and formation of the
20 lithium intercalated compound might be influenced by undesirable effects. Accordingly, the graphite after the pulverizing process has a great deal of crystalline structural defects. In a case when the above graphite is used as an active material for the negative electrode of the
25 lithium secondary battery, a disadvantage is caused that a large capacity can not be obtained. Furthermore, preferable performances of rapid charge and discharge are not obtained

because the lithium intercalation-deintercalation reaction is disturbed by the above defects.

Summary of the Invention

5 (1) Objects of the Invention:

The object of the present invention is to solve the above problems involved in the prior arts, to disclose a carbon material having a large lithium intercalation-deintercalation capacity and a method for manufacturing the same, and to provide a non-aqueous secondary battery which has
10 a large capacity and is superior in the rapid charging and discharging characteristics using the above disclosed materials.

(2) Methods of solving the Problems:

15 The crystalline structure of the graphite powder relating to the present invention has a feature that an existing fraction of the rhombohedral structure in the crystalline structure of the graphite is small (equal to or less than 20 %). Another feature is that an existing
20 fraction of the hexagonal structure is great (at least 80 %). The above existing fractions of the rhombohedral structure and the hexagonal structure can be determined by analyzing the intensity ratio of the peaks in the X-ray diffraction.

25 The graphite powder relating to the present invention is manufactured by a method comprising the steps of graphitizing treatment (heating at least 2000 °C) of raw

material such as oil cokes and coal cokes, pulverizing the graphitized raw material to powder, sieving the powder for obtaining the maximum particle diameter equal to or less than 100 μm , heating the powder at least 900 $^{\circ}\text{C}$ as a heat treatment, and further heating the powder at least 2700 $^{\circ}\text{C}$ for eliminating impurities such as Si. For instance, when the powder is heated at least 2700 $^{\circ}\text{C}$, Si, which is a main component of impurities, can be reduced to less than 10 ppm. The heat treatment of the powder for eliminating impurities can be omitted depending on the content of the impurities in the raw material. In the pulverizing process, various conventional pulverizers can be used. However, a jet mill is preferable, because pulverization with the jet mill generates the minimum destruction of the graphite crystalline structure in the raw material.

Furthermore, the graphite powder relating to the present invention can be obtained by immersing into an acidic solution containing at least one compound selected from a group consisted of sulfuric acid, nitric acid, perchloric acid, phosphoric acid, and fluoric acid as an immersing treatment after pulverizing the raw graphite to obtain graphite powder having a particle diameter equal to or less than 100 μm , subsequently washing with water, neutralizing, and drying.

The non-aqueous secondary battery for achieving the object of the present invention can be manufactured by using the graphite powder relating to the present invention as the

negative electrode active material, and the positive electrode is desirably composed of a material comprising a compound expressed by a chemical formula of Li_xMO_2 (where; X is in a range from zero to 1, and M is at least any one of chemical elements selected from a group of Co, Ni, Mn and Fe), or LiMn_2O_4 , that is a lithium transient metal complex oxide.

The active materials for the battery are generally used in a form of powder in order to proceed the charging and discharging reaction by increasing the surface area of the active material, which is a reaction field of the charging and discharging reaction. Therefore, the smaller the particle size of the powder is, the more will performance of the battery be improved. Furthermore, when the electrode is manufactured by applying an agent mixed with the active material and a binding agent to a current collector, the particle diameter of the active material is desirably equal to or less than 100 μm in view of applicability and maintaining preciseness of thickness of the electrode.

As for the negative electrode active material for the lithium secondary battery, natural graphite, artificial graphite, and others are disclosed. However, the above described reason, it is necessary to pulverize these materials. Therefore, in the pulverizing process, various graphite powder having a diameter equal to or less than 100 μm were prepared with various pulverizing methods using a ball mill, a jet mill, a colloidal mill and other apparatus, for various time. And, lithium intercalation-deintercalation

capacity of the various graphite powder were determined for searching a superior material for the negative electrode material of the lithium secondary battery.

However, the graphite powder obtained by the above method had the lithium intercalation-deintercalation amounts per weight in a range of 200-250 mAh/g, and their capacities as the material for the negative electrode of the lithium secondary battery were not enough.

In order to investigate the reason for the small capacity, crystalline structures of the above various graphite were determined by an X-ray diffraction method. FIG. 1 indicates an example of the results. Four peaks can be observed in a range of the diffraction angle (2θ , θ : Bragg angle) from 40 degrees to 50 degrees in the X-ray diffraction pattern. The peaks at approximately 42.3 degrees and 44.4 degrees are diffraction patterns of (100) plane and (101) plane of hexagonal structure of the graphite, respectively. The peaks at approximately 43.3 degrees and 46.0 degrees are diffraction patterns of (101) plane and (102) plane of rhombohedral structure of the graphite, respectively. As explained above, it was apparent that there were two kinds of crystalline structure in the pulverized graphite.

Further, the existing fraction (X) of the rhombohedral structure in the graphite powder was calculated by the following equation (Equation 1) based on the data of the observed peak intensity (P_1) of the (100) plane of the

hexagonal structure, the observed peak intensity (P_2) of the (101) plane of the rhombohedral structure, and a theoretical relationship of the intensity ratio in the X-ray pattern of graphite. As the result, it was revealed that the graphite
 5 having the rhombohedral structure was contained by approximately 30 % in all the graphite pulverized equal to or less than 100 μm in particle diameter.

$$X = 3P_2 / (11P_1 + 3P_2) \quad \dots (\text{Equation 1})$$

10

Similarly, the existing fraction (X) of the rhombohedral structure of the graphite powder was verified by the relationship of the observed peak intensity (P_1) of the (100) plane of the hexagonal structure, the observed
 15 peak intensity (P_3) of the (102) plane of the rhombohedral structure, and the theoretical relationship of the intensity ratio in the X-ray pattern of graphite. In this case, the following equation 2 was used instead of the equation 1. As the result, it was confirmed that the graphite having the
 20 rhombohedral structure was contained by approximately 30 % in all the graphite pulverized equal to or less than 100 μm in particle diameter.

$$X = P_3 / (3P_1 + P_3) \quad \dots (\text{Equation 2})$$

25

The reason for existing the two kinds of crystalline structure is assumed that the graphite itself has a

lubricating property, and the original graphite having the hexagonal structure transforms to the graphite having the rhombohedral structure by the pulverizing process with strong shocks. The graphite powder having a few microns in particle diameter obtained by further continued pulverization had a significantly broadened X-ray diffraction peak (P_4) at the (101) plane of the hexagonal structure, and it was revealed that the content of amorphous carbon in the graphite was increased because the half band width of the peak was increased. Accordingly, the reason for the small lithium intercalation-deintercalation capacity of the conventional graphite powder can be assumed that the crystalline structure of the graphite has transformed to the rhombohedral structure and generated the amorphous carbon, and proceeding of the lithium intercalation-deintercalation reaction is disturbed by the rhombohedral structure and the amorphous carbon.

Analysis on the impurities of the graphite powder revealed that the impurities such as Si, Fe, and others were contained more than 1000 ppm. Naturally, in addition to the impurities contained in the raw material, impurities from processing apparatus such as a ball mill, a jet mill, and the like can be mixed into the graphite at the pulverizing process. Therefore, the influence of the above impurities can be assumed as another reason for the small capacity in addition to the above formation of the rhombohedral structure and amorphous carbon.

In accordance with the present invention, the graphite powder having a particle diameter equal to or less than 100 μm , wherein the content of the above described rhombohedral structure is less than 30 % and the content of the amorphous carbon is small, has been developed. Additionally, the content of Si in particular, which is the main component of the impurities in the graphite powder, has been decreased to equal to or less than 10 ppm. Therefore, extremely high purity is one of the features of the graphite relating to the present invention. The particle diameter equal to or less than 100 μm is determined with an intention to use the graphite for the battery, as described previously. Therefore, when the graphite of the present invention is used for other usages, the particle diameter of the graphite is not necessarily restricted to equal to or less than 100 μm .

Hereinafter, details of the graphite powder relating to the present invention, and the method for manufacturing the same is explained.

Two methods (manufacturing method 1 and manufacturing method 2) for obtaining the graphite having a small fraction of the rhombohedral structure are disclosed.

(Manufacturing method 1)

As for raw material (raw graphite) for the graphite powder of the present invention, both natural graphite and artificial graphite can be used. In particular, flaky natural graphite is preferable. Among the above raw graphite, the one of which maximum diffraction peak in the

X-ray diffraction pattern by the $\text{CuK}\alpha$ line is appeared at a diffraction angle (2θ , θ : Bragg angle) in a range from 26.2 degrees to 26.5 degrees, that is, an interval between two graphite layers is equal to or less than 0.34 nm, is
5 desirable. Because, the graphite powder containing a small amount of the rhombohedral structure can be obtained from the high crystalline raw material.

As for the pulverizing apparatus for crushing the raw graphite to the particle diameter equal to or less than 100
10 μm , a jet mill is desirable. The reason is that the amorphous carbon is generated less with the jet mill than the case when another pulverizing apparatus is used.

The pulverized raw graphite (raw powder) contains the graphite having the rhombohedral structure by approximately
15 30 % as previously described. Then, in accordance with the present manufacturing method 1, the existing fraction of the rhombohedral structure is made decrease by the following heat treatment.

The heat treatment is performed at least 900 $^{\circ}\text{C}$ under an
20 inert gas atmosphere. As for the inert gas, nitrogen gas, argon gas, and the like is used. The inert gas atmosphere can also be maintained by covering the raw powder with cokes to seal it from the atmosphere.

The heat treatment is the most important process in the
25 present invention for transforming the rhombohedral structure to the hexagonal structure. It is necessary to perform the heat treatment after pulverization of the raw

graphite (more preferably, at the last stage of the graphite powder manufacturing process of the present invention).

If the heat treatment is performed before the pulverization of the graphite and subsequently the graphite is pulverized, the graphite powder containing the rhombohedral structure as small as possible, which is the object of the present invention, can not be obtained. The graphite powder containing the rhombohedral structure graphite as small as possible can be obtained only by the heat treatment after the pulverizing process (more preferably, at the last stage of the graphite powder manufacturing process of the present invention) as the present invention.

The raw graphite powder contains Al, Ca, Fe, and particularly much of Si, as impurities. The impurities can be eliminated by heating and sublimating the materials at least 2700 °C. Therefore, the heating temperature in the heat treatment is preferably at least 2700 °C in order to perform a purification treatment concurrently.

(Manufacturing method 2)

The raw graphite and the pulverizing process is as same as the above manufacturing method 1.

The graphite powder of the present invention can be obtained by treating the graphite powder obtained by the pulverizing process with an acidic solution containing at least one compound selected from a group consisted of sulfuric acid, nitric acid, perchloric acid, phosphoric acid, and fluoric acid, and subsequently washing with water,

neutralizing, and drying. During the treatment, a compound is formed with anions in the above acidic solution and the graphite, and the rhombohedral structure graphite is eliminated by the formation of the compound. The anions from the acidic solution in the compound are eliminated from the compound during the washing, the neutralizing, and the drying, and the graphite powder relating to the present invention can be obtained.

The crystalline structure of the graphite powder of the present invention obtained by the above manufacturing methods 1 and 2 was analyzed by an X-ray diffraction. The ratio of the P_1 and P_2 , (P_2/P_1) , was less than 0.92, and the half band width of the P_4 was less 0.45 degrees. The ratio of the P_1 and P_3 , (P_3/P_1) , was less than 0.75.

By substituting the above observed data for the equations 1 and 2, the fact that the existing fraction of the rhombohedral structure has been decreased less than 20 % and the existing fraction of the hexagonal structure has been increased at least 80 % was confirmed. Simultaneously, the content of Si was confirmed to be less than 10 ppm from the result of impurity analysis.

Then, an electrode was prepared using the graphite powder of the present invention as an active material, and lithium intercalation-deintercalation capacity was studied. As the result, the lithium intercalation-deintercalation capacity of the graphite powder of the present invention was 320-360 mAh/g per unit weight of the active material, and

the capacity was significantly improved in comparison with the capacity of the conventional graphite material (200-250 mAh/g). Furthermore, it was found that the preferable existing fraction of the rhombohedral structure was equal to 5 or less than 10 %, because the less the existing fraction of the rhombohedral structure in the graphite powder of the present invention is, the more will the capacity be increased.

Accordingly, the rhombohedral structure is evidently a 10 crystalline structure which hardly intercalate or deintercalate lithium. Therefore, it is assumed that the high lithium intercalation-deintercalation capacity of the graphite powder of the present invention is achieved by especially decreasing the existing fraction of the 15 rhombohedral structure and increasing the existing fraction of the hexagonal structure.

The feature of the lithium secondary battery of the present invention is in using the graphite powder of the present invention as the negative active material. The 20 lithium secondary battery relating to the present invention has a large load capacity, and a high energy density can be realized.

As the result of an evaluation on the characteristics of the lithium secondary battery of the present invention, it 25 was confirmed that the lithium secondary battery of the present invention had a superior performance in rapid charging and discharging characteristics, and decreasing

ratio of the capacity was improved at least 30 % in comparison with the conventional lithium battery under a same rapid charging and discharging condition. The reason for the improvement can be assumed that the reversibility
5 for the lithium intercalation-deintercalation reaction of the graphite of the present invention is improved in comparison with the conventional carbon material by decreasing the existing fraction of the rhombohedral structure and eliminating the influence of the impurities
10 such as Si.

As the positive active material for the lithium secondary battery of the present invention, materials such as Li_xCoO_2 , Li_xNiO_2 , $\text{Li}_x\text{Mn}_2\text{O}_4$, (where, X is in a range 0-1) and the like are desirable because a high discharge voltage of at least
15 3.5 V can be obtained, and the reversibility of the charge and discharge of the positive electrode itself is superior.

As for the electrolytic solution, a mixed solvent composed of ethylene carbonate mixed with any one selected from a group consisted of dimethoxyethane, diethylcarbonate, dimethylcarbonate, methylethylcarbonate, γ -butyrolactone, methyl propionate, and ethyl propionate, and at least one of
20 electrolytes selected from a group consisted of salts containing lithium such as LiClO_4 , LiPF_6 , LiBF_4 , LiCF_3SO_3 , and the like are used. It is desirable to adjust the lithium
25 concentration in a range 0.5- 2 mol/l, because, an electric conductivity of the electrolytic solution is favorably large.

Brief Description of Drawings

Figure 1 indicates an X-ray diffraction pattern of the conventional graphite,

Figure 2 indicates an X-ray diffraction pattern of the graphite powder relating to the embodiment 1 of the present invention (heat treatment temperature: 900 °C),

Figure 3 indicates an X-ray diffraction pattern of the graphite powder relating to the embodiment 1 of the present invention (heat treatment temperature: 2850 °C),

Figure 4 indicates an X-ray diffraction pattern of the graphite powder prepared in the comparative example 1,

Figure 5 indicates an X-ray diffraction pattern of the graphite powder relating to the embodiment 2 of the present invention,

Figure 6 indicates a schematic cross section of the battery used in the embodiment 3 and the comparative example 2,

figure 7 is a graph indicating a relationship between the electrode potential and the lithium intercalation-deintercalation capacity,

Figure 8 is a partial cross section of the lithium secondary battery prepared in the embodiment 5 of the present invention,

Figure 9 is a graph indicating a relationship between the discharge capacity and the number of repeating the charge and the discharge cycles,

Figure 10 is a graph indicating a relationship between

the discharge capacity and the charging and discharging current,

Detailed Description of the Embodiments

5

Referring to drawings, embodiments of the present invention are explained hereinafter.

Embodiment 1

10 Flaky natural graphite which was produced from Madagascar was used as the raw material, and the raw material was pulverized to be powder, of which particle diameter was equal to or less than 46 μm , by a jet mill. The powder was sieved to obtain raw material powder. The average
15 diameter of the raw material powder was 8.0 μm .

Subsequently, the raw material powder was processed with a heat treatment by heating at 900 $^{\circ}\text{C}$ or 2850 $^{\circ}\text{C}$ for ten days under a nitrogen atmosphere, and the graphite powder of the present invention was obtained.

20 The crystalline structures of the graphite powder of the present invention and the raw material powder were analyzed by an X-ray diffraction method using an apparatus RU-200 made by Rigaku Denki, and the impurity content was analyzed by an inductively coupled plasma spectrometry (ICP) using an
25 apparatus P-5200 made by Hitachi.

The X-ray diffraction patterns of the graphite powder of the present invention, which have been observed under a

condition of X-ray tube voltage; 40 kV, X-ray tube current; 150 mA, and X-ray source; CuK α line, are shown in FIGs. 2 and 3. FIG. 2 is the pattern obtained by the heat treatment at 900 °C, and FIG. 3 is the pattern obtained by the heat treatment at 2850 °C. The X-ray diffraction patterns of the graphite powder of the present invention in both FIG 2 and FIG. 3 indicate that the peaks at diffraction angles of 43.3 degrees and 46.0 degrees, both of which belong with the rhombohedral structure, are decreased by either of the above heat treatments.

The amount of Si contained in the graphite powder of the present invention as an impurity was 1140 ppm when the heating temperature was 900 °C, and 27 ppm when the heating temperature was 2850 °C. Therefore, it is revealed that a highly purified graphite powder, of which Si is eliminated, can be obtained by heat treatment at a high temperature at least 2700 °C, by which Si can be eliminated.

Comparative example 1

In order to compare with the embodiment of the present invention, the non-pulverized raw graphite was heated at 2850 °C, and subsequently pulverized to obtain the graphite powder. The X-ray pattern of the graphite powder obtained by the above process is shown in FIG. 4. It is apparent from FIG. 4 that the peaks at diffraction angles of 43.3 degrees and 46.0 degrees, both of which belong with the rhombohedral structure, are not decreased. That means, the rhombohedral

structure can not be eliminated by the above process.

Embodiment 2

In accordance with the embodiment 2, the raw graphite
5 was pulverized by a jet mill to less than 100 μm in particle
diameter. Then, the graphite powder was immersed into a
mixed acid of sulfuric acid and nitric acid for a whole day.
Subsequently, washing with distilled water and
neutralization with a dilute aqueous solution of sodium
10 hydroxide were performed. The graphite powder obtained by
the above process was dried at 120 $^{\circ}\text{C}$ to obtain the graphite
powder of the present invention. The X-ray pattern of the
graphite powder obtained by the above process is shown in
FIG. 5. The peaks at diffraction angles of 43.3 degrees and
15 46.0 degrees, both of which belong with the rhombohedral
structure, are decreased. Accordingly, it was found that the
rhombohedral structure was eliminated by the above process.

Embodiment 3

20 In accordance with the embodiment 3, a carbon electrode
was prepared using the graphite powder of the present
invention as an electrode active material, and the lithium
intercalation-deintercalation capacity, in other words, a
load capacity of the negative electrode in the lithium
25 secondary battery was studied with the electrode.

Mixed agents slurry were prepared by mixing 90 % by
weight in total solid of the graphite powder of the present

invention prepared in the embodiment 1, 10 % by weight of polyvinylidene fluoride (PVDF) as a binder, and N-methyl-2-pyrrolidone of which heating temperature were 900 °C and 2850°C, respectively. The mixed agents slurry was applied on
5 a plane of a sheet of copper foil of 10 µm thick, and dried in vacuum at 120 °C for one hour. After the vacuum drying, an electrode was fabricated by roller pressing, of which thickness was in a range 85-90 µm. The average amount of the applied mixed agents per unit area was 10 mg/cm². The
10 electrode was prepared by cutting the copper foil applied with the mixed agents into a sheet of 10 mm X 10 mm.

FIG. 6 is a schematic cross section of a battery used for studying the lithium intercalation-deintercalation capacity of the present electrode. The battery has a structure,
15 wherein a working electrode current collector 30, the electrode of the present invention 31, which is a working electrode, a separator 32, a lithium metal 33, which is a counter electrode, a counter electrode current collector 34 are piled and inserted into a battery vessel 35, and a battery
20 lid 36 is screwed for fixing. A reference electrode made of lithium metal 37 is attached to the battery. As for the electrolytic solution, a mixed solvent of ethylene carbonate and diethylcarbonate by 1:1 in volume and lithium hexafluorophosphate were used with lithium concentration of
25 1 mol/l.

The intercalation-deintercalation of lithium was repeated by applying a constant current between the working

electrode and the counter electrode, and the capacity was determined. The terminated potentials of the intercalation and the deintercalation of the working electrode were set as 0 V and 0.5 V, respectively.

5

Comparative example 2

In order to compare with the embodiment of the present invention, a carbon electrode was prepared with the graphite powder obtained in the comparative example 1 by the same
10 method as the embodiment 3, and the load capacity (the amount of lithium intercalation-deintercalation) was determined. The same study was performed on the electrode prepared with the conventional graphite powder (the same powder as the raw powder in the embodiment 1).

15 A result of comparison on the lithium intercalation-deintercalation behavior of the electrode in the embodiment 3 (the present invention) with the electrode in the comparative example 2 (prior art) and the electrode prepared with the conventional graphite powder is explained
20 hereinafter. FIG. 7 is a graph indicating a relationship between the lithium intercalation-deintercalation capacity and an electrode potential at the fifth cycle, wherein the capacity becomes stable, after repeating the intercalation-deintercalation of lithium. In FIG. 7, the curve
25 indicates the potential variation of the electrode prepared with the graphite powder, of which heating temperature at the heat treatment was 900 °C, in the embodiment 3. The curve

41 indicates the potential variation of the electrode prepared with the graphite powder, of which heating temperature at the heat treatment is 2850 °C, in the embodiment 3. The curve 42 indicates the potential variation of the electrode prepared with the conventional graphite powder, and the curve 43 indicates the potential variation of the electrode prepared with the graphite powder which has been prepared in the comparative example 1 by the reversely ordered processes. The intercalation capacity and the deintercalation capacity for lithium in both the cases of using the conventional graphite in the comparative example 2 (the curve 42) and the graphite in the comparative example 1 (the curve 43) were less than 250 mAh/g per unit weight of the active materials. On the contrary, in the case of the embodiment 3 (the curves 40, 41), wherein the graphite powder prepared in the embodiment 1 was used as the active material, both the intercalation capacity and the deintercalation capacity for lithium were more than 300 mAh/g per unit weight of the active materials. That means, a large load capacity was obtained by using the graphite powder having a small existing fraction of the rhombohedral structure relating to the present invention. Furthermore, the case (the curve 41) using the graphite powder highly purified by heating up to 2850 °C indicates the largest values in both the intercalation capacity and the deintercalation capacity for lithium in FIG. 7.

Embodiment 4

The embodiment 4 was performed in order to confirm the influence of treating time in the heat treatment of the present invention. In the embodiment 4, the graphite powder of the present invention was obtained in accordance with the substantially same manner as the embodiment 1 (under a nitrogen atmosphere, the raw powder was heated at 2850 °C). However, the treating time of the heat treatment was varied in a range from 0 hours to 30 days.

10 The existing fraction of the rhombohedral structure was determined from the peak intensity in X-ray diffraction patterns. Furthermore, as same as the embodiment 3, the electrodes were prepared with the obtained graphite powders, and the intercalation-deintercalation reactions of lithium
15 were repeatedly performed. The result on the lithium intercalation-deintercalation capacity at the fifth cycle is shown in Table 1.

Table 1

	Heating time	The existing fraction of the rhombohedral structure (%)	Lithium intercalation capacity (mAh/g)	Lithium deintercala- tion capacity (mAh/g)
5	0 hours	27.3	249	235
	4 hours	18.2	332	320
	10 hours	14.6	345	325
	1 day	13.8	343	334
	3 days	11.3	355	338
	5 days	9.7	368	351
	10 days	7.1	365	360
10	30 days	3.9	366	361

In accordance with the above result, it is apparent that the smaller the existing fraction of the rhombohedral structure is, the more will the lithium intercalation-deintercalation capacity be increased. In particular, the existing fraction equal to or less than 10 % is desirable.

Embodiment 5

The present embodiment uses a cylindrical lithium secondary battery. A fundamental structure of the secondary battery is shown in FIG. 8. In FIG.8, the member assigned with the numeral mark 50 is a positive electrode. Similarly, a negative electrode 51, a separator 52, a positive electrode tab 53, a negative electrode tab 54, a positive electrode lid 55, a battery vessel 56, and a gasket 57 are shown.

The lithium secondary battery shown in FIG. 8 was prepared by the following steps. Mixed positive electrode agents slurry was prepared by mixing 88 % by weight in total solid of LiCoO_2 as an active material for the positive electrode, 7 % by weight of acetylene black as a conductive agent, 5 % by weight of polyvinylidene fluoride (PVDF) as a binder, and N-methyl-2-pyrrolidone.

Similarly, mixed negative electrode agents slurry was prepared by mixing 90 % by weight in total solid of the graphite powder of the present invention as an active material for the negative electrode, 10 % by weight of polyvinylidene fluoride (PVDF) as a binder, and N-methyl-2-pyrrolidone.

The mixed positive electrode agents slurry was applied onto both planes of a sheet of aluminum foil of 25 μm thick, and dried in vacuum at 120 $^{\circ}\text{C}$ for one hour. After the vacuum drying, an electrode of 195 μm thick was fabricated by roller pressing. The average amount of the applied mixed agents per unit area was 55 mg/cm^2 . The positive electrode was prepared by cutting the aluminum foil applied with the mixed agents into a sheet of 40 mm in width and 285 mm in length. However, portions of 10 mm in length from both ends of the positive electrode were not applied with the mixed agents for the positive electrode, the aluminum foil was bared, and one of the bared portion was welded to the positive electrode tab by ultrasonic bonding.

The mixed negative electrode agents slurry was applied onto both planes of a sheet of copper foil of 10 μm thick, and dried in vacuum at 120 $^{\circ}\text{C}$ for one hour. After the vacuum drying, an electrode of 175 μm thick was fabricated by roller pressing. The average amount of the applied mixed agents per unit area was 25 mg/cm^2 . The negative electrode was prepared by cutting the copper foil applied with the mixed agents into a sheet of 40 mm in width and 290 mm in length. However, as same as the positive electrode, portions of 10 mm in length from both ends of the negative electrode were not applied with the mixed agents for the negative electrode, the copper foil was bared, and one of the bared portion was welded to the negative electrode tab by ultrasonic bonding.

A fine pored film made of polypropylene of 25 μm thick and 44 mm in width was used as a separator. The positive electrode, the separator, the negative, and the separator were piled in the order described above, and the pile was rolled to form a bundle of the electrodes. The bundle was contained in a battery vessel, the negative electrode tab was welded to the bottom of the battery vessel, and a drawn portion for caulking the positive electrode lid was fabricated. An electrolytic solution prepared by adding lithium hexafluorophosphate by 1 mol/l into a mixed solvent containing ethylene carbonate and diethylcarbonate by 1:1 in volume was filled the battery vessel, the positive electrode tab was welded to the positive electrode lid, and the positive electrode lid was caulked to the battery vessel to form the battery.

Using the battery which had been prepared by the above steps, the charge and discharge were repeated under a condition that the charging and discharging current was 300 mA, and respective of the terminated potentials of the charge and the discharge was 4.2 V and 2.8 V. Furthermore, the charging and the discharging current was varied in a range from 300 mA to 900 mA, and the rapid charge and rapid discharge were performed.

Comparative example 3

In order to compare with the present invention, a lithium secondary battery was manufactured by the method as

same as the embodiment 5 using the conventional graphite powder (the raw powder for the graphite powder of the present invention), and the battery characteristics was determined as same as the embodiment 5.

5 The result of comparison on the characteristics of the lithium secondary battery of the embodiment 5 (the present invention) and the comparative example 3 (prior art) is explained hereinafter.

FIG. 9 indicates variation in discharge capacity of the
10 lithium secondary battery when the charge and discharge of the battery were repeated. The curve 60 indicates the discharge capacity of the embodiment 5. The curve 61 indicates the discharge capacity of the comparative example 3. In the embodiment 5, the maximum discharge capacity was 683 mAh,
15 and a ratio in the discharge capacity after 200 cycles to the maximum capacity was 86 %. While, in the comparative example 3, the maximum discharge capacity was 492 mAh, and a ratio in the discharge capacity after 200 cycles to the maximum capacity was 63 %.

20 FIG. 10 indicates a relationship between the charging and discharging current and the discharge capacity when the rapid charge and rapid discharge were performed. The curve 70 indicates the discharge capacity of the embodiment 5. The curve 71 indicates the discharge capacity of the comparative
25 example 3. With the charging and discharging current of 900 mA, the discharge capacity of the embodiment 5 was 573 mAh, while the discharge capacity of the comparative example 3

was 256 mAh. The ratio of decreasing the discharge capacity in the respective of the present cases to the discharge capacity in the case of charging and discharging current of 300 mAh/g were 16 % and 48 %, respectively. Therefore, in accordance with using the graphite powder of the present invention as the active material for the negative electrode, the decreasing ratio of the capacity was improved by at least 30 %, and it became apparent that the lithium secondary battery relating to the present invention had an excellent characteristics in rapid charge and discharge.

Embodiment 6

Mixed positive electrode agents slurry was prepared using LiMn_2O_4 as a positive electrode active material, and the positive electrode was prepared by applying the mixed positive electrode agents slurry onto both planes of a sheet of aluminum foil by as same as the embodiment 5. The average amount of the applied mixed agents per unit area was 65 mg/cm^2 , and the electrode thickness after fabrication by roller pressing was $230 \text{ }\mu\text{m}$. The positive electrode was prepared by cutting the aluminum foil applied with the mixed agents into a sheet of 40 mm in width and 240 mm in length. However, portions of 10 mm in length from both ends of the positive electrode were not applied with the mixed agents for the positive electrode. The negative electrode was as same as the negative electrode prepared in the embodiment 5. Then, the lithium secondary battery of the present embodiment

was prepared by the same method as the embodiment 5 such as forming an electrodes bundle, inserting the electrodes bundle into a vessel, welding bottom of the vessel, adding an electrolytic solution, caulking a positive electrode lid, and others.

Using the battery, charge and discharge were repeated under a condition of charging and discharging current of 300 mA, and the terminated potential of the charge and discharge of 4.2 V and 2.8 V, respectively. As the result, the maximum discharge capacity was 581 mAh, and a ratio in the discharge capacity after repeating the charging and discharging reactions 200 cycles to the maximum discharge capacity was 84 %. The above result indicates that the charging and discharging characteristics of the present embodiment is superior to the comparative example 3.

A lithium secondary battery which has a high energy density and an excellent charging and discharging characteristics can be obtained by using the graphite powder, which is superior in reversibility of the intercalation-deintercalation reaction of lithium, of which maximum particle size is less than 100 μm , wherein the existing fraction of the rhombohedral structure in the crystalline structure is less than 20 %, as the active material for the negative electrode of the battery.

WHAT IS CLAIMED IS:

1. A method for manufacturing graphite powder, comprising the steps of:
 - graphitizing raw graphite by heating raw graphite to at least 2000°C, to produce graphitized raw graphite;
 - pulverizing said graphitized raw graphite, to produce pulverized graphite;
 - sieving said pulverized graphite for obtaining graphite powder having a maximum particle diameter of 100 μm ;
 - heating said graphite powder as a heat treatment for transforming the crystalline structure to hexagonal structure; and
 - further heating said graphite powder, at a higher temperature than said heat treatment for transforming the crystalline structure, for eliminating impurities.
2. A method for manufacturing graphite powder as claimed in claim 1, wherein
 - the temperature of said heat treatment for transforming crystalline structure to hexagonal structure is in a range from 900°C to 1100°C.
3. A method for manufacturing graphite powder as claimed in claim 1, wherein
 - the temperature of said heat treatment for eliminating impurities is in a range from 2700°C to 2900°C.

4. A method for manufacturing graphite powder as claimed in claim 1, wherein

the heat treatments are performed after said pulverizing process.

5. A method for manufacturing graphite powder, comprising the steps of:

graphitizing raw graphite by heating raw graphite at least 2000°C, to produce graphitized raw graphite;

pulverizing said graphitized raw graphite, to produce pulverized graphite;

sieving said pulverized graphite for obtaining graphite powder having a maximum particle diameter of 100 μm ;

immersing said graphite powder into an acidic solution as an immersing treatment;

washing with water;

neutralizing; and

drying.

6. A method for manufacturing graphite powder as claimed in claim 5, wherein

said acidic solution contains at least one compound selected from a group consisting of sulfuric acid, nitric acid, perchloric acid, phosphoric acid, and fluoric acid.

7. A method for manufacturing graphite powder as claimed in any one of claims 1-5, wherein

said pulverizing is performed by a jet-mill.

8. A method for manufacturing graphite powder as claimed in any one of claims 1-6, wherein

said raw graphite has a diffraction angle for the maximum diffraction peak in a range from 26.2 degrees to 26.5 degrees in an X-ray diffraction pattern with the CuK α line.

9. A method for manufacturing graphite powder as claimed in claim 7, wherein

said raw graphite has a diffraction angle for the maximum diffraction peak in a range from 26.2 degrees to 26.5 degrees in an X-ray diffraction pattern with the CuK α line.

10. a non-aqueous secondary battery comprising:

a positive electrode,
a negative electrode, and

electrolytic solution, which is charged or discharged by repeating a reaction of intercalating and deintercalating ions at said positive electrode and said negative electrode, respectively, wherein

said graphite powder composing said negative electrode has a particle size equal to or smaller than 100 μm , and

said negative electrode comprises graphite powder having a fraction of a rhombohedral structure equal to or less than 20% by weight.

11. A non-aqueous secondary battery as claimed in claim 10, wherein

said graphite powder has a fraction of a hexagonal structure equal to or more than 80% by weight.

12. A non-aqueous secondary battery comprising:
a positive electrode,
a negative electrode, and
electrolytic solution, which is charged or discharged by repeating a reaction of intercalating and deintercalating ions at said positive electrode and said negative electrode, respectively, wherein
said graphite powder composing said negative electrode has a particle size equal to or smaller than 100 μm ,
and
said negative electrode comprises graphite powder having a fraction of a rhombohedral structure equal to or less than 10% by weight.

13. A non-aqueous secondary battery as claimed in claim 12, wherein said graphite powder has a fraction of a hexagonal structure equal to or more than 90% by weight.

14. A non-aqueous secondary battery comprising:
a positive electrode,
a negative electrode, and
electrolytic solution, which is charged or discharged by repeating a reaction of intercalating and deintercalating ions at said positive electrode and said negative electrode, respectively, wherein

said negative electrode comprises graphite powder having a particle size equal to or smaller than 100 μm ,

said graphite powder has both a hexagonal structure and a rhombohedral structure, and

said graphite powder has a fraction of the rhombohedral structure equal to or less than 20% by weight, and a fraction of the hexagonal structure equal to or more than 80% by weight.

15. A non-aqueous secondary battery, made by a method comprising the steps of:

laminating graphite electrodes with a lithium group oxide; and

enclosing said graphite electrodes into a cell vessel with an electrolyte solution, wherein

said graphite electrodes are manufactured by the steps of:

granulating the graphite to graphite powder having a particle size equal to or smaller than 100 μm ,

treating said graphite powder by heating at 900°C or higher, after said granulating, and

fabricating said graphite electrodes by subjecting the heat-treated graphite powder to pressing.

16. A non-aqueous secondary battery according to claim 15, wherein said treating said graphite powder by heating is performed so as to modify crystallinity of the graphite powder such that a fraction of the graphite powder having

rhombohedral structure is equal to or less than 20 % by weight.

17. A non-aqueous secondary battery according to claim 16, wherein, in said treating said graphite powder by heating, said crystallinity of the graphite powder is modified so that a fraction of the graphite powder having hexagonal structure is equal to or greater than 80% by weight.

18. A non-aqueous secondary battery according to claim 16, wherein crystallinity of the graphite powder is modified during the heat treatment so that a fraction of the graphite powder having rhombohedral structure is equal to or less than 10% by weight.

19. A non-aqueous secondary battery, made by a method comprising the steps of:

laminating graphite electrodes with a lithium group oxide; and

enclosing said graphite electrodes into a cell vessel with an electrolyte solution, wherein

said graphite electrodes are manufactured by the steps of:

granulating the graphite to graphite powder having a particle size equal to or smaller than 100 μ m,

immersing said graphite powder into an acidic solution as an immersing treatment, said acidic solution containing at least one compound selected from a group

consisting of sulfuric acid, nitric acid, perchloric acid, phosphoric acid and fluoric acid, and then washing said graphite powder with water, neutralizing, and drying said graphite powder, and

fabricating said graphite electrodes by subjecting the dried graphite powder to pressing.

20. A method of manufacturing a lithium secondary battery, comprising the steps of:

laminating graphite electrodes with a lithium group oxide; and

enclosing said graphite electrodes into a cell vessel with an electrolyte solution, wherein

said graphite electrodes are manufactured by the steps of:

granulating the graphite to graphite powder having a particle size equal to or smaller than 100 μm ,

treating said graphite powder by heating at 900°C or higher, after said granulating, and

fabricating said graphite electrodes by subjecting the heat-treated graphite powder to pressing.

21. A method of manufacturing a lithium secondary battery, comprising the steps of:

laminating graphite electrodes with a lithium group oxide; and

enclosing said graphite electrodes into a cell vessel with an electrolyte solution, wherein

said graphite electrodes are manufactured by the steps of:

granulating the graphite to graphite powder having a particle size equal to or smaller than 100 μm ,

immersing said graphite powder into an acidic solution as an immersing treatment, washing said graphite powder, neutralizing said graphite powder, and drying said graphite powder, and

fabricating said graphite electrodes by subjecting the dried graphite powder to pressing.

22. A method of manufacturing a lithium secondary battery according to claim 21, wherein said acidic solution contains at least one compound selected from a group consisting of sulfuric acid, nitric acid, perchloric acid, phosphoric acid and fluoric acid.

Abstract of the Disclosure

Objects of the present invention is to provide a carbon material having a superior reversibility in lithium intercalation-deintercalation reaction, and a non-aqueous secondary battery using the carbon material as an active material for a negative electrode, which has a high energy density and an excellent rapid charging and discharging characteristics.

Graphite powder having a maximum particle diameter of less than 100 μm and an existing fraction of rhombohedral structure in the crystalline structure of less than 20 % is used as an active material for the negative electrode of the non-aqueous secondary battery. The graphite powder can be obtained by pulverizing raw graphite with a jet mill, and subsequently treating the powder at a temperature equal to or higher than 900 $^{\circ}\text{C}$.

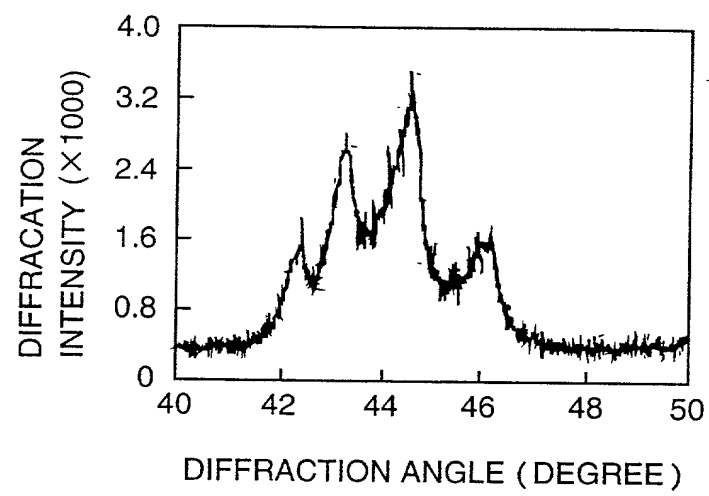
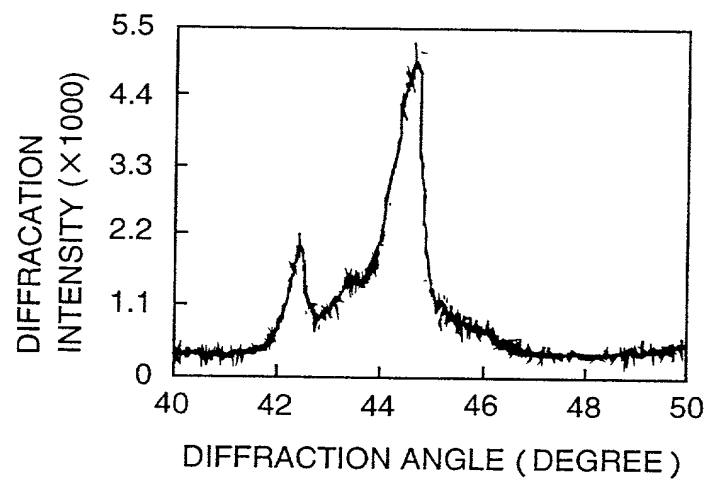
FIG .1*FIG .2*

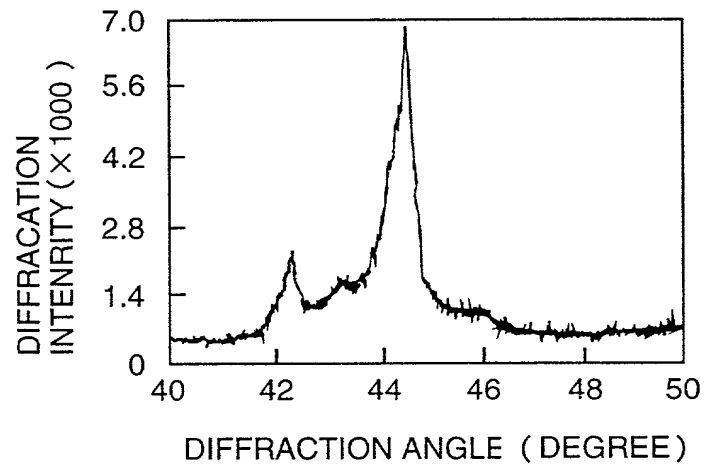
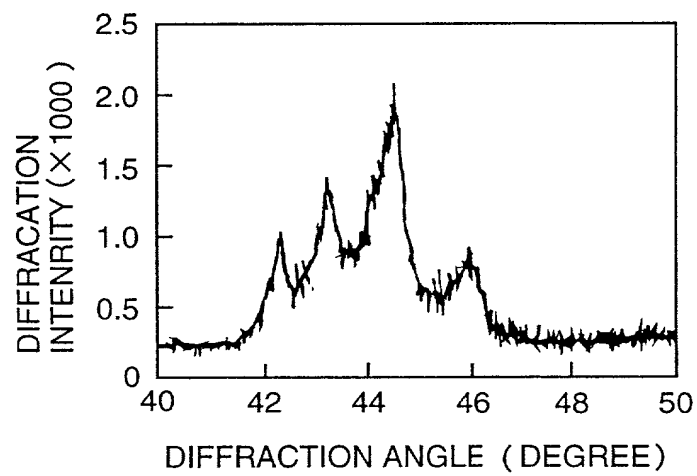
FIG .3*FIG .4*

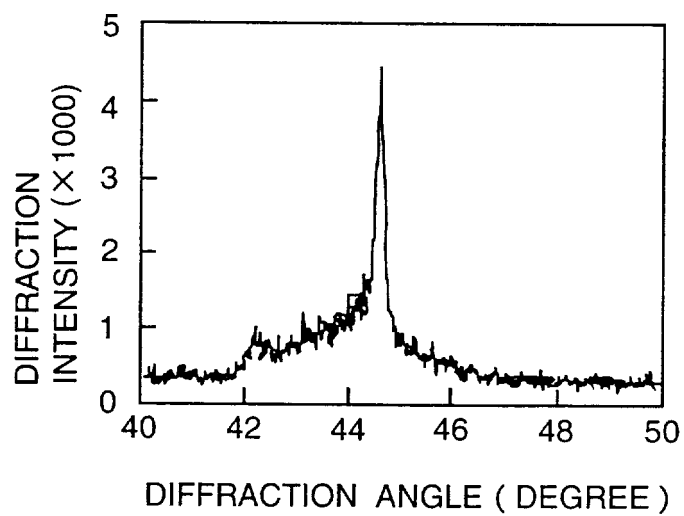
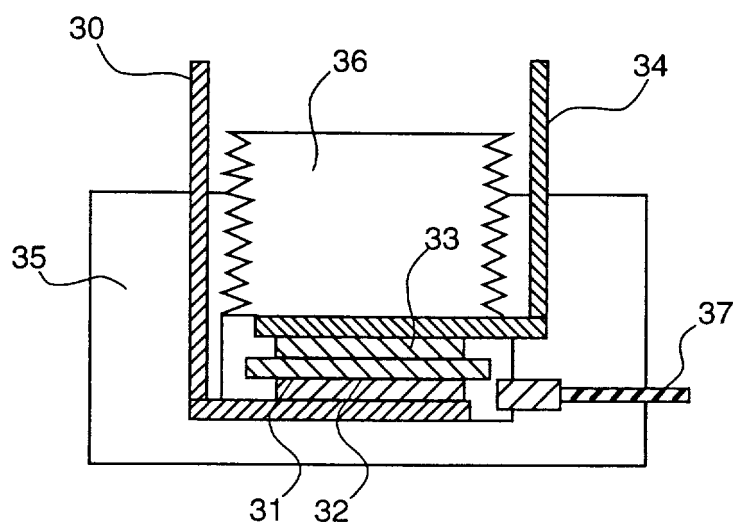
FIG .5*FIG .6*

FIG .8

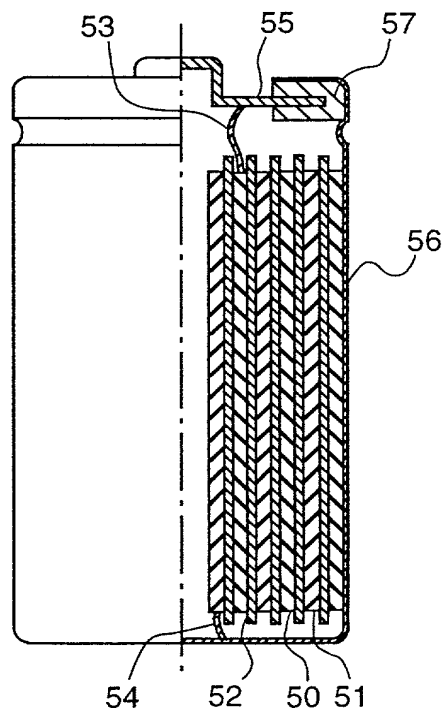
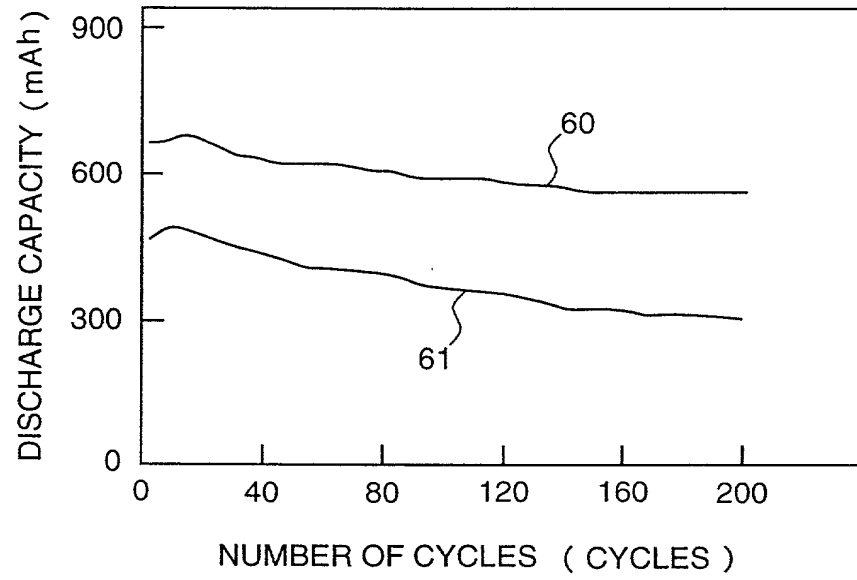
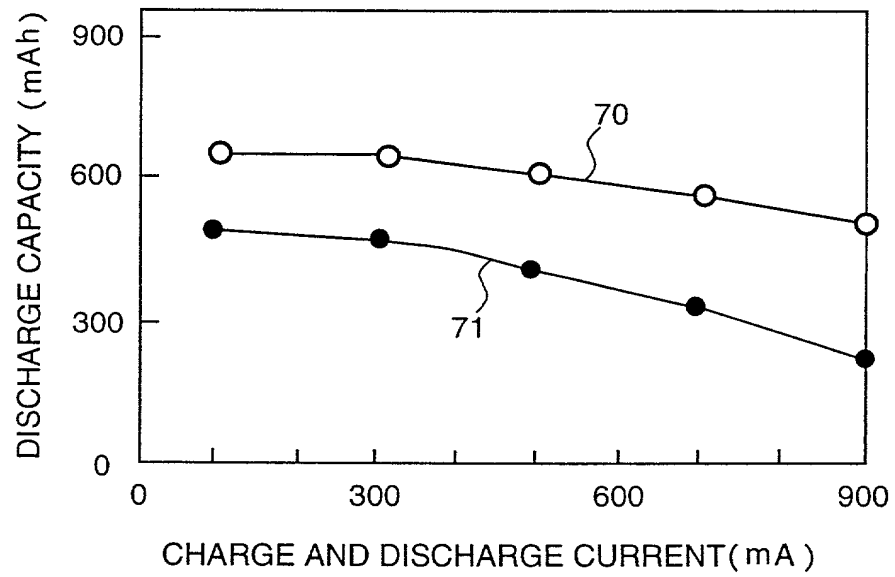


FIG .9*FIG .10*

DECLARATION AND POWER OF ATTORNEY FOR PATENT APPLICATION

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name, I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled:

NON-AQUEOUS SECONDARY BATTERY AND A MANUFACTURING METHOD OF GRAPHITE POWDER

the specification of which (check one)

☐

is attached hereto.

☒

was filed on April 10, 1996

as Application Serial No. 08/630,501

and was amended on _____

(if applicable)

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to the examination of this application in accordance with Title 37, Code of Federal Regulations, §1.56(a).

I, hereby claim foreign priority benefits under Title 35, United States Code, § 119 of any foreign application(s) for patent or inventor's certificate listed below and have also identified below any foreign application for patent or inventor's certificate having a filing date before that of the application on which priority is claimed:

Prior Foreign Application(s)

Priority Claimed

<u>7-84196</u> (Number)	<u>Japan</u> (Country)	<u>10/04/1995</u> (Day/Month/Year Filed)	<input checked="" type="checkbox"/> Yes	<input type="checkbox"/> No
_____ (Number)	_____ (Country)	_____ (Day/Month/Year Filed)	<input type="checkbox"/> Yes	<input type="checkbox"/> No
_____ (Number)	_____ (Country)	_____ (Day/Month/Year Filed)	<input type="checkbox"/> Yes	<input type="checkbox"/> No
_____ (Number)	_____ (Country)	_____ (Day/Month/Year Filed)	<input type="checkbox"/> Yes	<input type="checkbox"/> No
_____ (Number)	_____ (Country)	_____ (Day/Month/Year Filed)	<input type="checkbox"/> Yes	<input type="checkbox"/> No
_____ (Number)	_____ (Country)	_____ (Day/Month/Year Filed)	<input type="checkbox"/> Yes	<input type="checkbox"/> No

I hereby claim the benefit under Title 35, United States Code, § 120 of any United States application(s) listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States application in the manner provided by the first paragraph of Title 35, United States Code, § 112, I acknowledge the duty to disclose material information as defined in Title 37, Code of Federal Regulations, § 1.56(a) which occurred between the filing date of the prior application and the national or PCT international filing date of this application:

_____ (Application Serial No.)	_____ (Filing Date)	_____ (Status: patented, pending, abandoned)
_____ (Application Serial No.)	_____ (Filing Date)	_____ (Status: patented, pending, abandoned)
_____ (Application Serial No.)	_____ (Filing Date)	_____ (Status: patented, pending, abandoned)
_____ (Application Serial No.)	_____ (Filing Date)	_____ (Status: patented, pending, abandoned)

(Continued on Page 2)

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I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further, that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United State Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

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Date _____	Inventor _____	
Residence _____	Citizenship _____	
Post Office Address _____		
Date _____	Inventor _____	
Residence _____	Citizenship _____	
Post Office Address _____		